FIG. 2

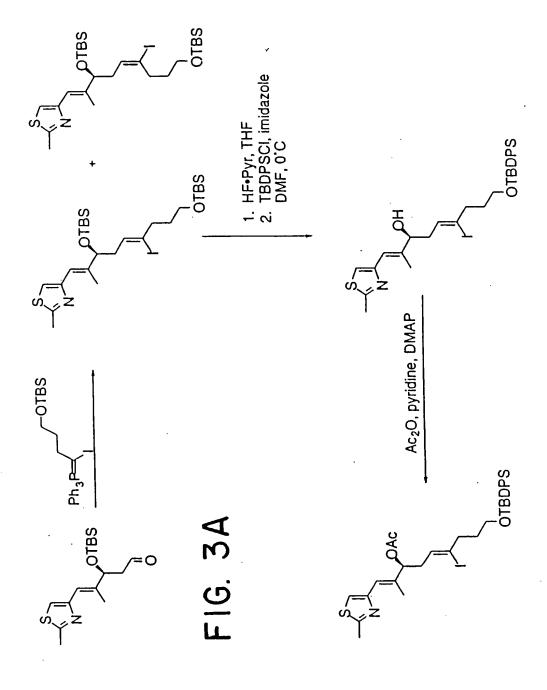
OAC
$$R = CH_2OTBS \\ (CH_2)_3OTBDPS$$

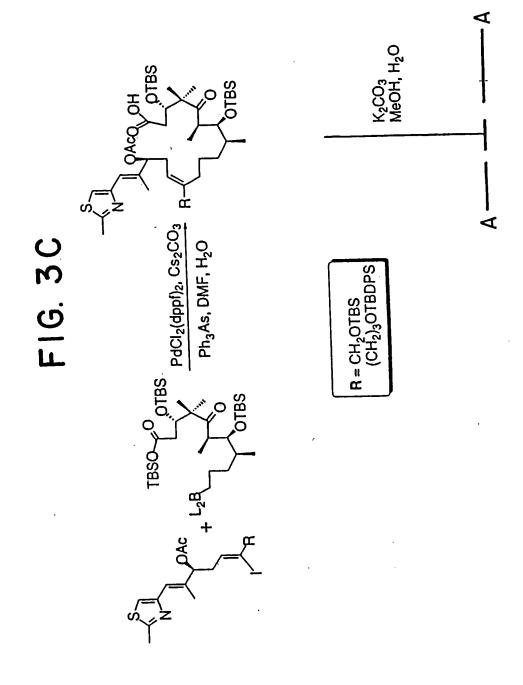
$$R = CH_2OTBS \\ (CH_2)_3OTBDPS$$

$$R = CH_2OH$$

(CH₂)₃OH

$$(CH_2)_3OBz$$
 CF_3
 CH_2O
 $N=N$
 CF_3
 CF_3
 CCF_3





F16.3E

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50% for three steps

*17 steps from known starting materials vs. 27 steps for aldol macrocyclization

cis epoxide (epothilone A)

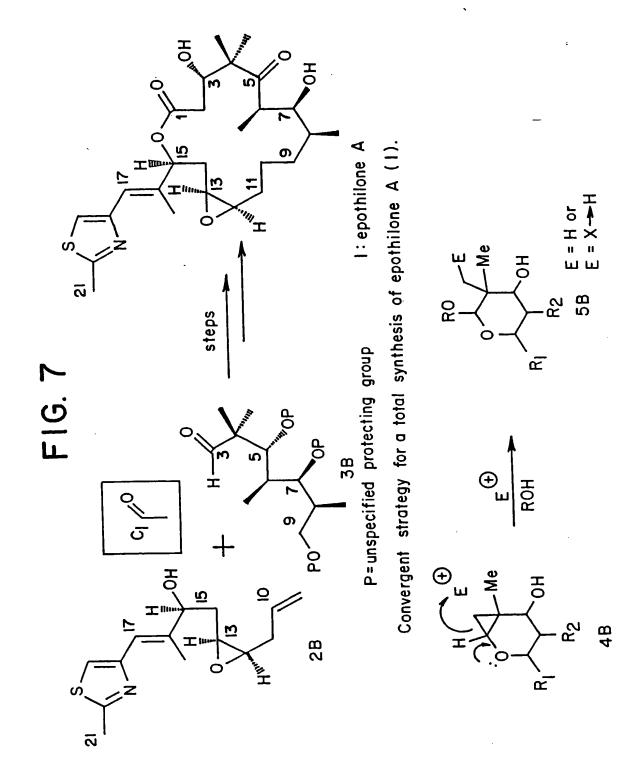


FIG. 10

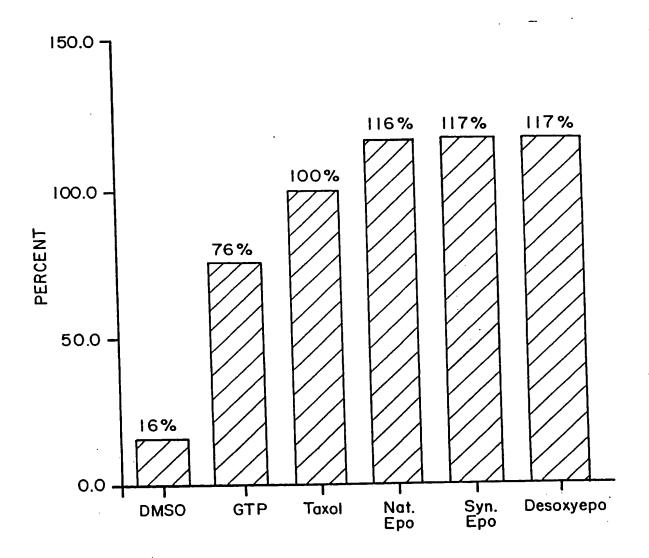


FIG. 11

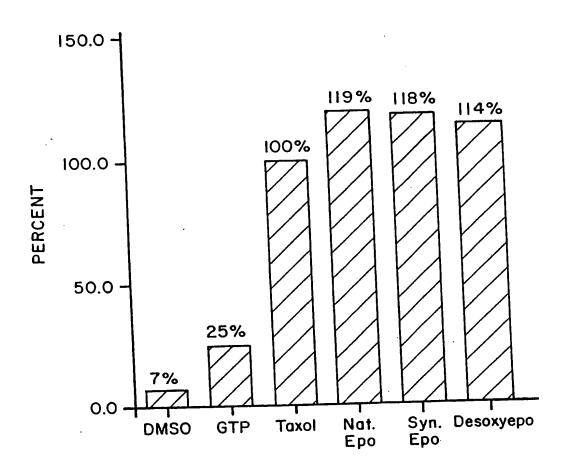
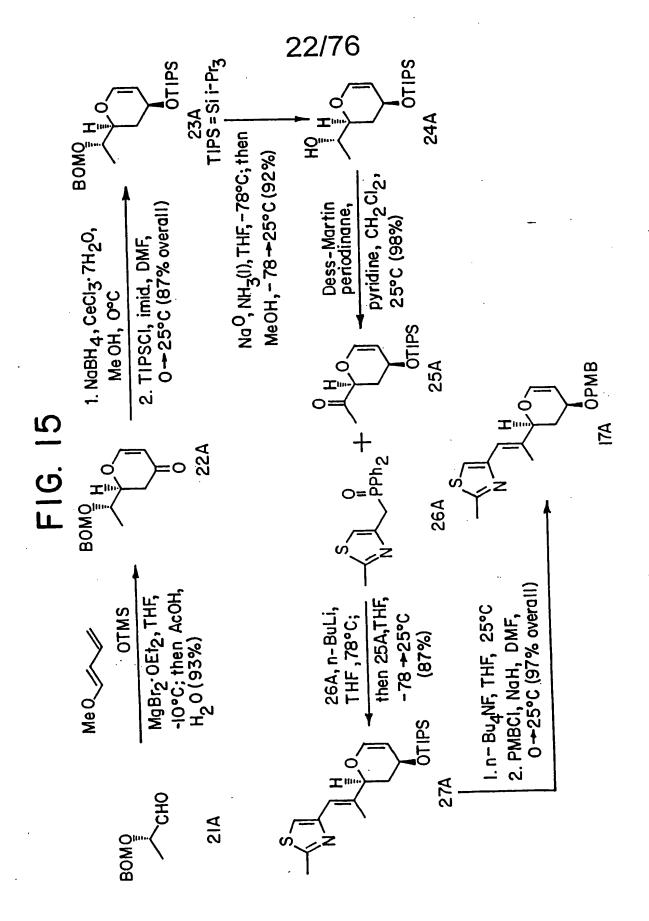
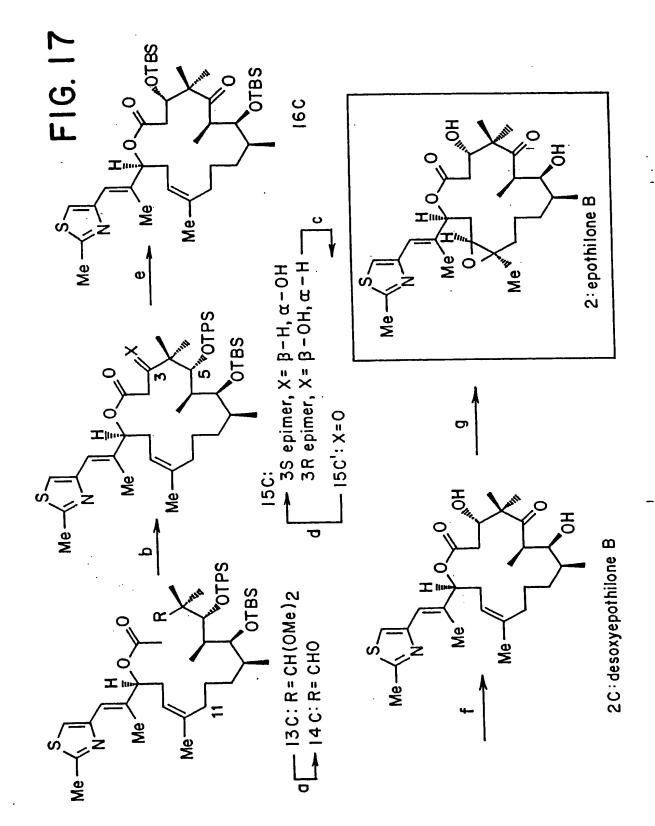


FIG. 14B





28/76 9 20 OTBS TBS = Sit-buMe2 80 - 9D#R=t-butyl ► 10D#R=TBS a,b *****0TBS FIG. 20B

FIG. 21

F1G. 22 A

FIG. 22B

19D TEA, 4-DMAP, C6H6 F1G. 24A FIG. 24B MeOH/H₂O 88% 18D minor product from suzuki coupling reaction HF-pyridine, THF 95%

FIG. 25A

87% ee

FIG. 25B

R=H is the only compound completed, F and CF₃ are nearly R = H, F, CF3 ∞mpleted FIG. 26B 20%

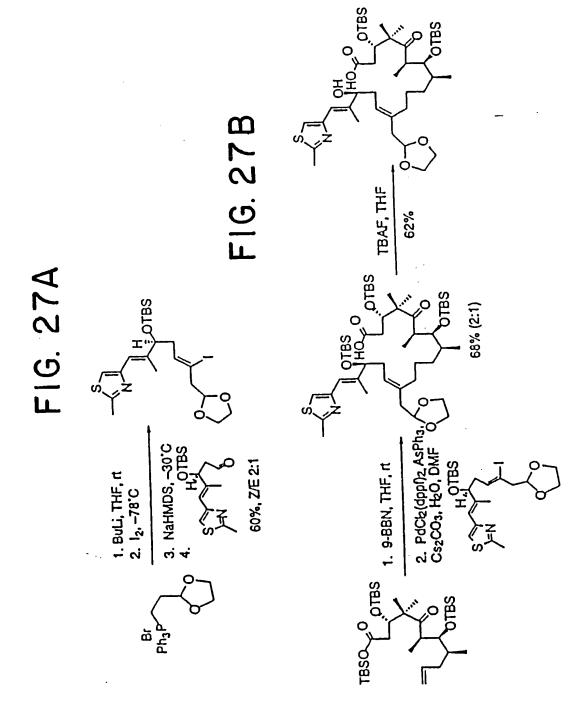


FIG. 28A

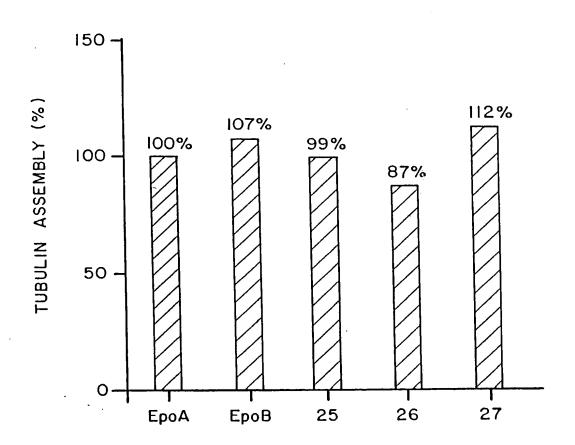
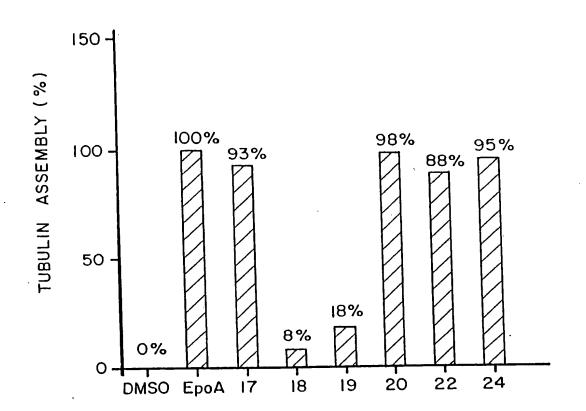
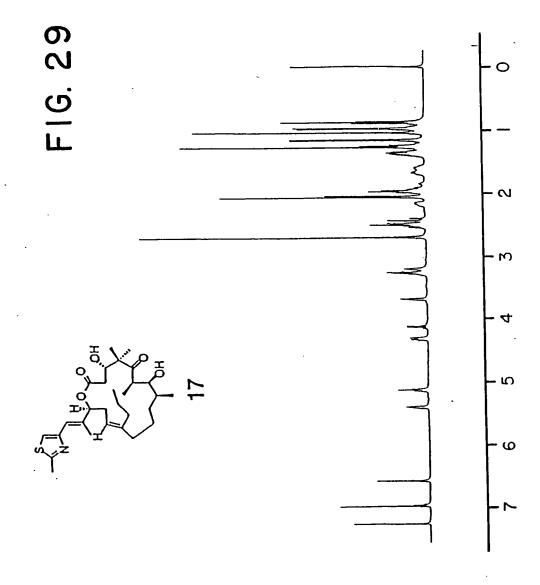
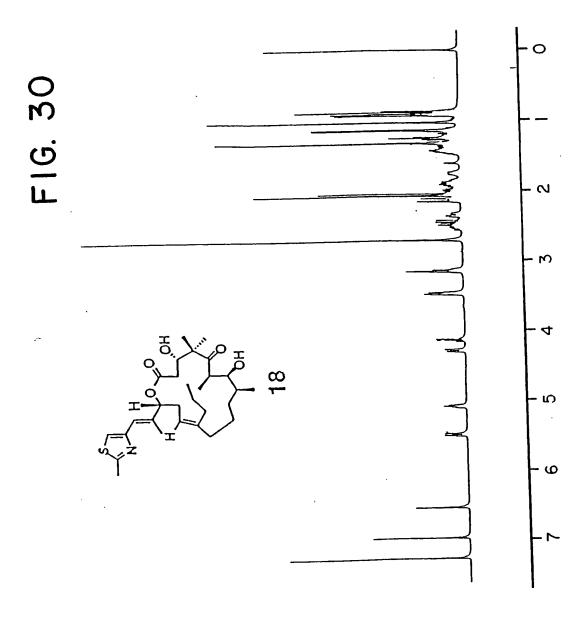
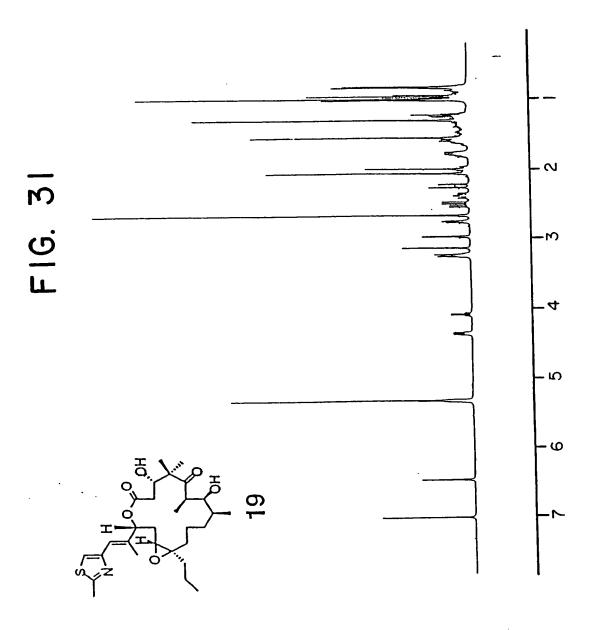


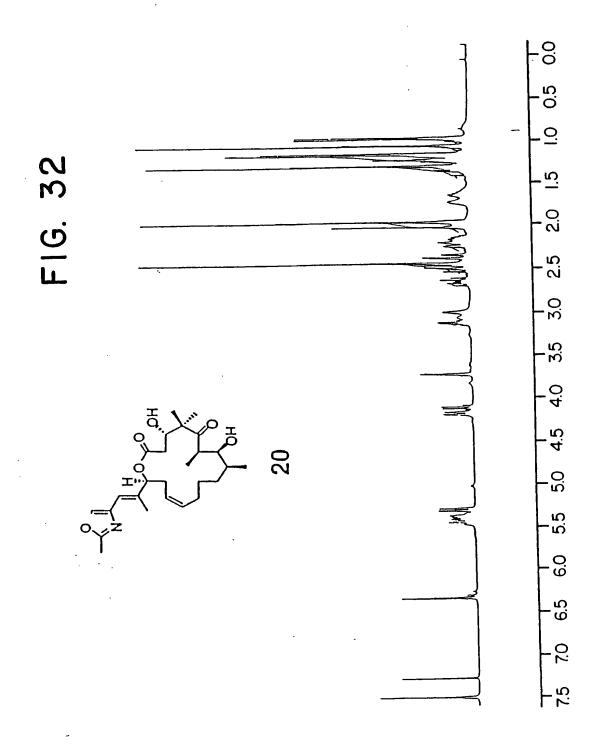
FIG. 28B

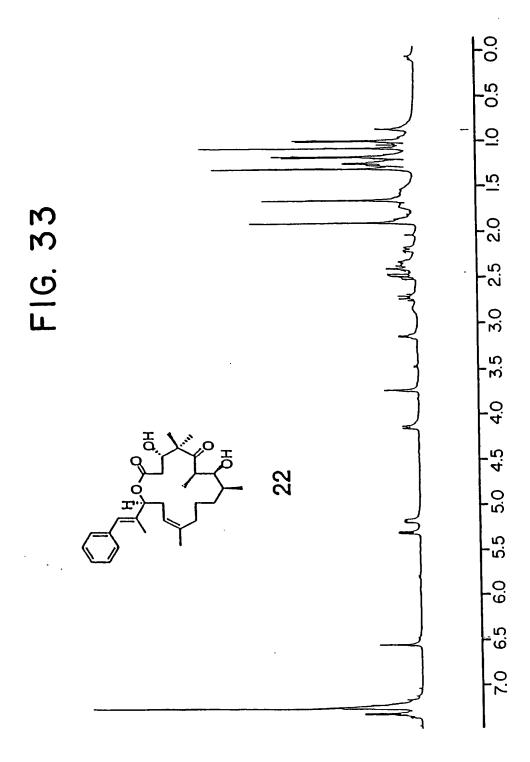


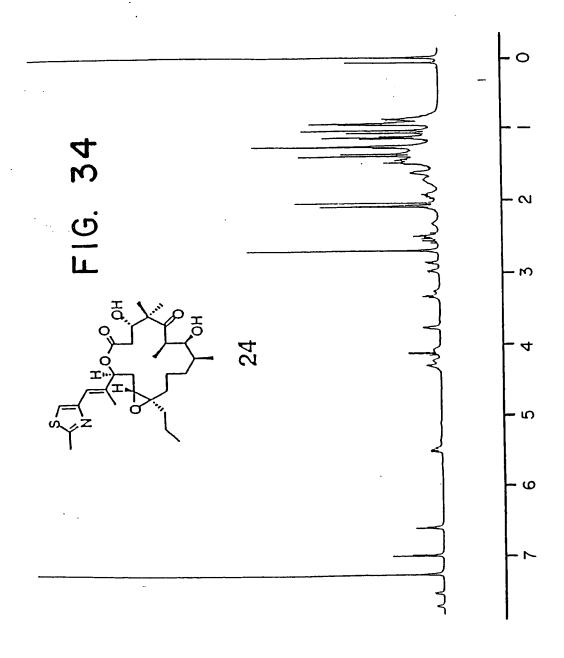


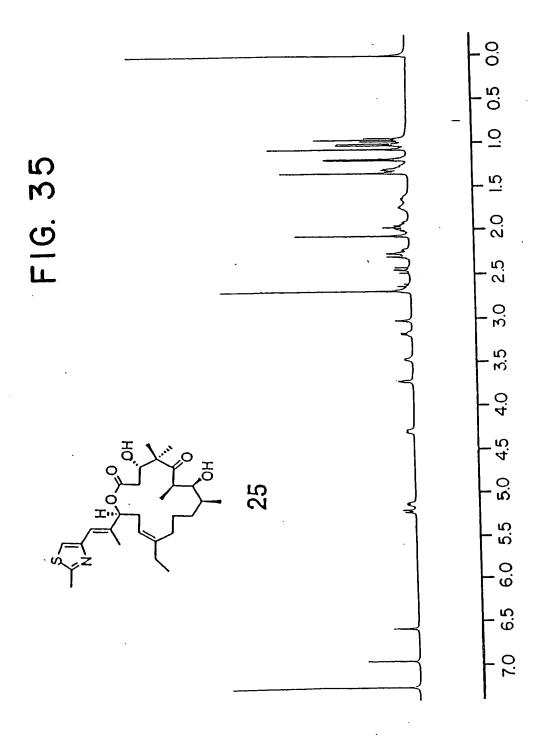


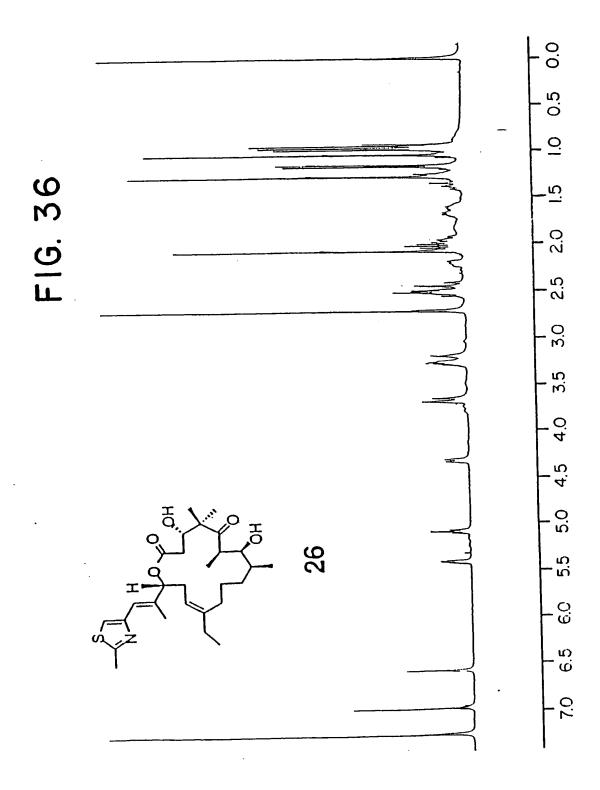


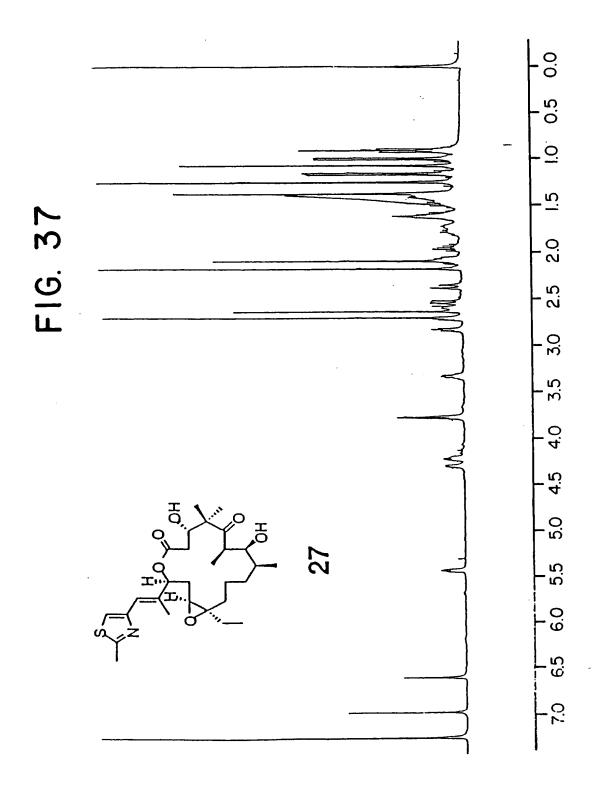












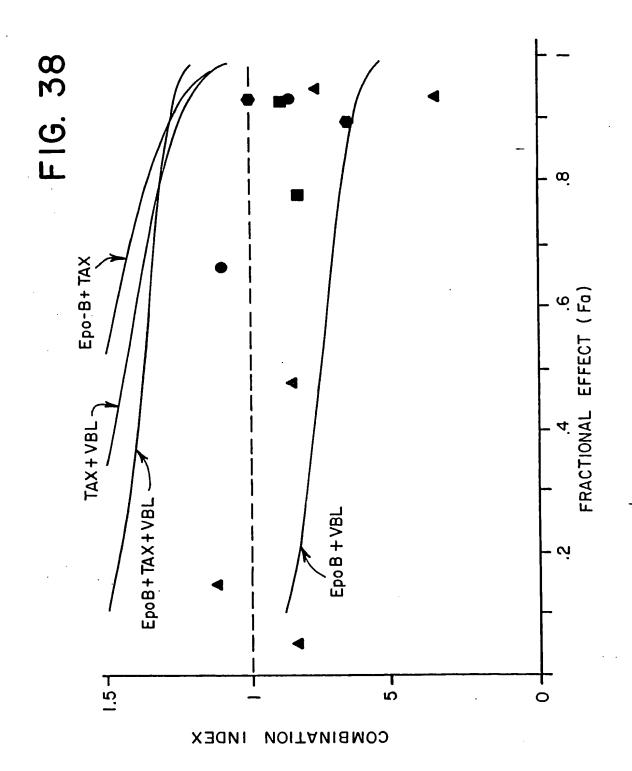


FIG. 39A

synthetic epothilone B

8 (0.00044) [0.0026]

FIG. 41A

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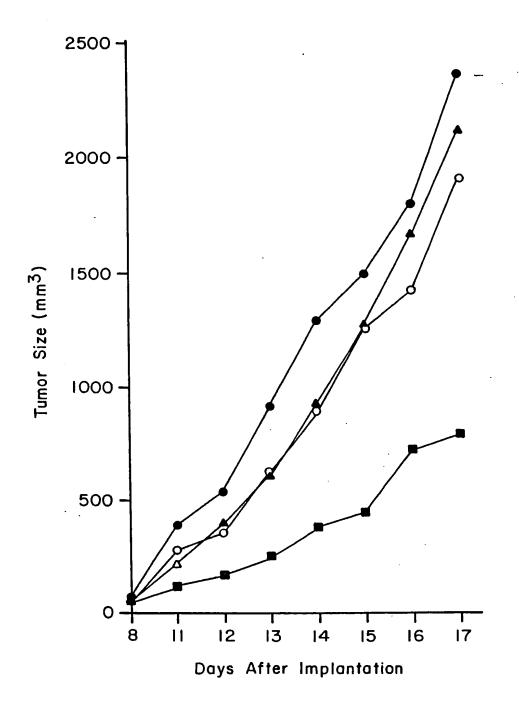
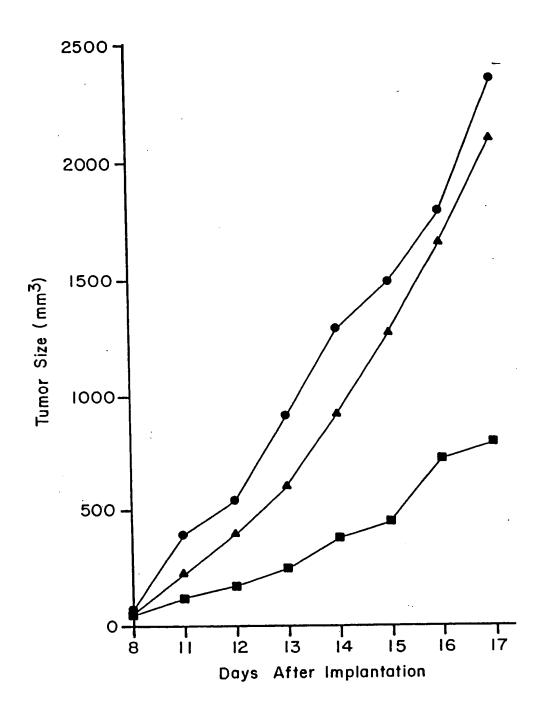
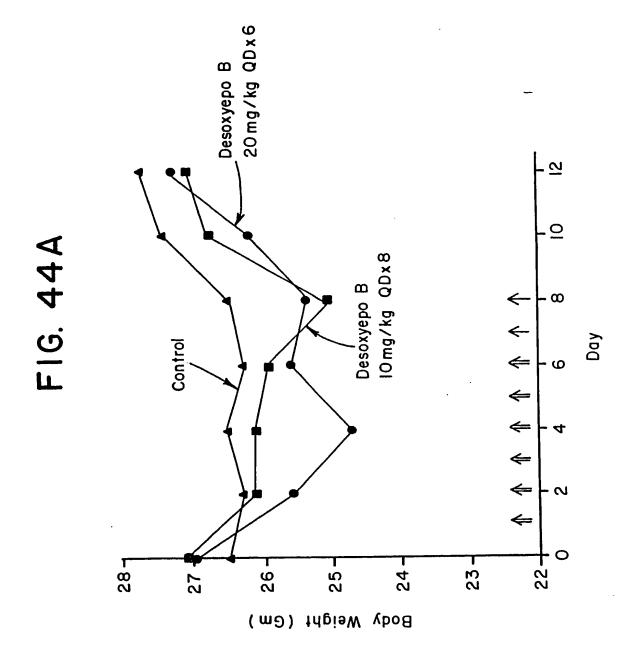
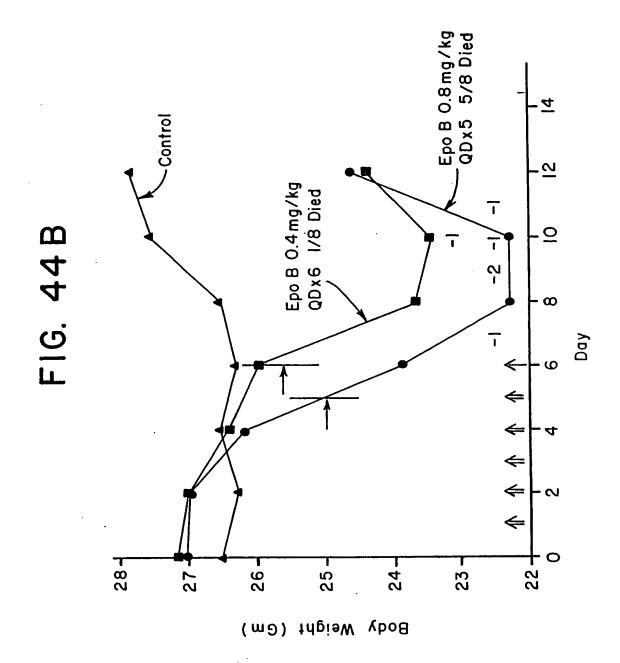


FIG. 43B

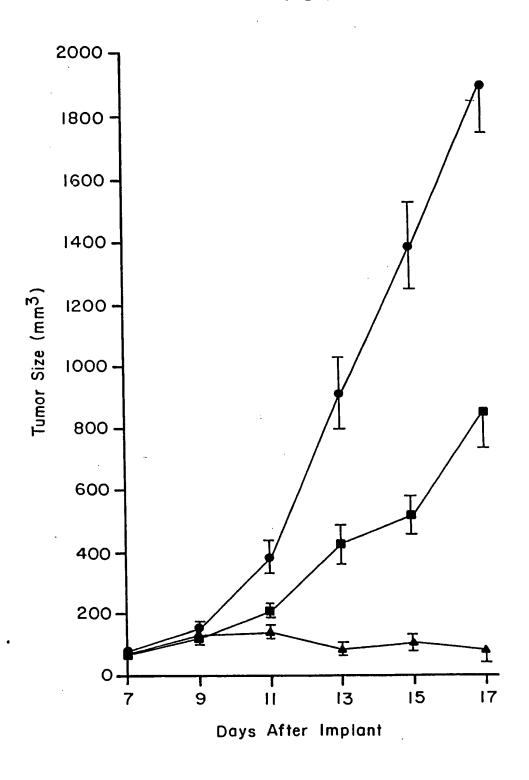






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FIG. 45A



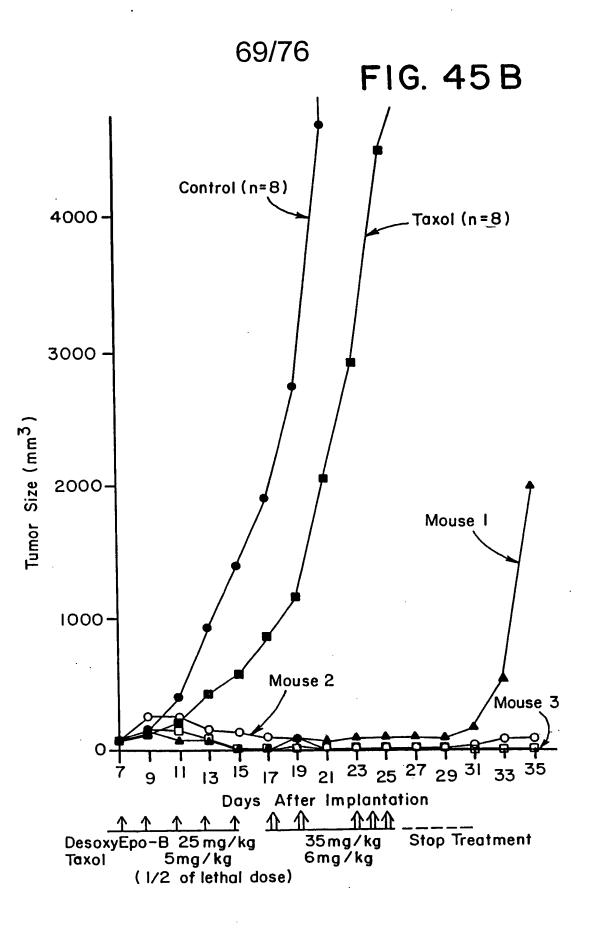


FIG. 46

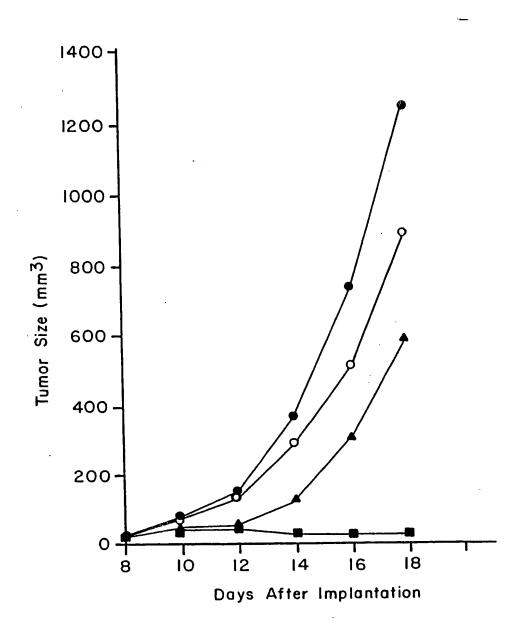


FIG. 47

